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AMENDMENTS TO THE SPECIFICATION:

Please amend the specification as follows:

Paragraph bridging pages 8-9, lines 36-37 and 1-10, respectively:

Amino acid and nucleotide sequences can also be evaluated either manually by one skilled in the art, or by using computer-based sequence comparison and identification tools that employ algorithms such as BLAST (Basic Local Alignment Search Tool; Altschul et al. (1993) *J. Mol. Biol.* 215:403-410[; see also [in situ hybridization of bacterial colonies or bacteriophage plaques\). In addition, short oligonucleotides of 12 or more nucleotides may be used as amplification primers in PCR in order to obtain a particular nucleic acid fragment comprising the primers.](http://www.ncbi.nlm.nih.gov/BLAST/)

Paragraph at page 21, lines 25-41:

cDNA clones encoding AMP or adenosine deaminases were identified by conducting BLAST (Basic Local Alignment Search Tool; Altschul et al. (1993) *J. Mol. Biol.* 215:403-410[; see also [Nat. Genet. 3:266-272\) provided by the NCBI. For convenience, the P-value \(probability\) of observing a match of a cDNA sequence to a sequence contained in the searched databases merely by chance as calculated by BLAST are reported herein as "pLog" values, which represent the negative of the logarithm of the reported P-value. Accordingly, the greater the pLog value, the greater the likelihood that the cDNA sequence and the BLAST "hit" represent homologous proteins.](http://www.ncbi.nlm.nih.gov/BLAST/)